LYCORIS RUBUS BRIGHTENING BB SPF40 PA PLUS PLUS- octinoxate, titanium dioxide, zinc oxide cream Lycoris Co., Ltd.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

ACTIVE INGREDIENT

Active Ingredients: Ethylhexyl methoxycinnamate 7.5%, Titanium dioxide 5%, Zinc oxide 5%

INACTIVE INGREDIENT

Inactive Ingredients: Water, Cyclopentasiloxane, Butylene glycol, Isoamyl p-methoxycinnamate, Cetyl PEG/PPG-10/1 Dimethicone, Betaine, Cetyl ethylhexanoate, Polysorbate 60, Dimethicone, Sodium chloride, Sodium hyaluronate, PEG-40 stearate, Iron oxides, Bentonite, Talc, Stearic acide, Phenoxyethanol, Methylparaben, Ethylparaben, Butylparaben, Propylparaben, Isobutylparaben, Tocopheryl acetate, Perfume, Rubus hirsutus fruit/leaf extract, Camellia Japonica Ovary Powder, Centella asiatica extract, Lavandula angustifolia (Lavender) extract, Glycyrrhiza glabra (Licorice) root extract, Triticum vulgare (Wheat) flour lipids, Allantoin, Lysine HCL, Disodium EDTA, Proline, Sodium ascorbyl phosphate, Linum usitatissimum (Linseed) seed extract, Hibiscus esculentus fruit extract, Aloe barbadensis leaf juice, Arctium lappa root extract, Hibiscus mutabilis flower extract, Corchorus olitorius leaf extract, Dimethyl sulfone

PURPOSE

Purpose: Sunscreen

WARNINGS

Warnings: 1. In case of having following symptoms after using this, you're advised to stop using it immediately. If you keep using it, the symptoms will get worse and need to consult a dermatologist. 1) In case of having problem such as red rash, swollenness, itching, stimulation during usage. 2) In case of having the same symptoms above on the part you put this product on by direct sunlight. 2. You are banned to use it on the part where you have a scar, eczema, or dermatitis. 3. In the case of getting it into your eyes, you have to wash it immediately.

KEEP OUT OF REACH OF CHILDREN

Keep out of reach of babies and children

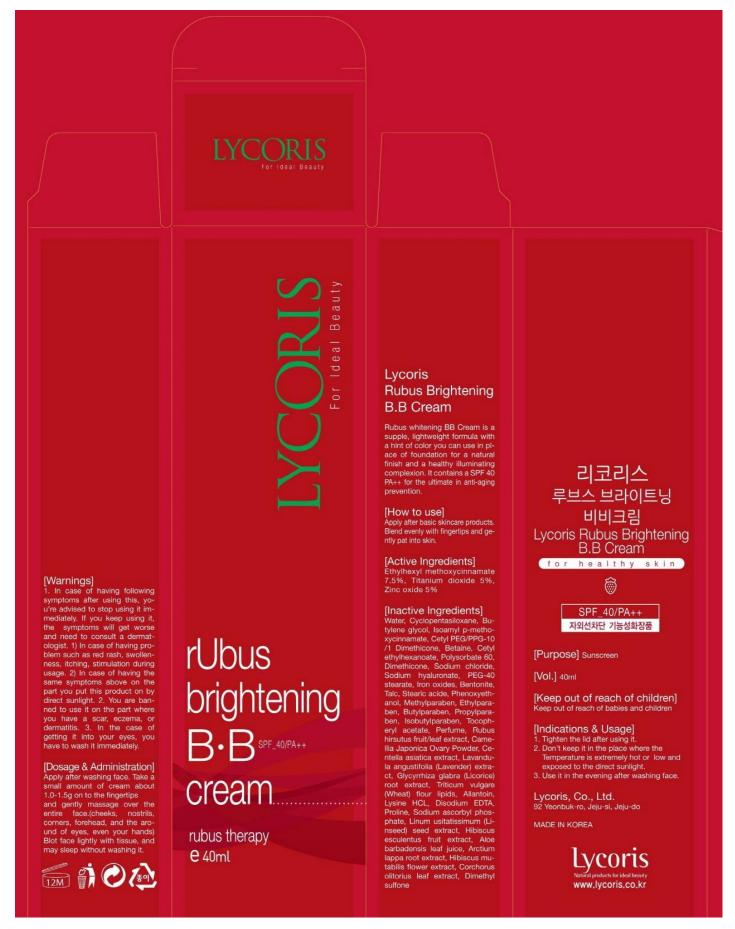
INDICATIONS AND USAGE

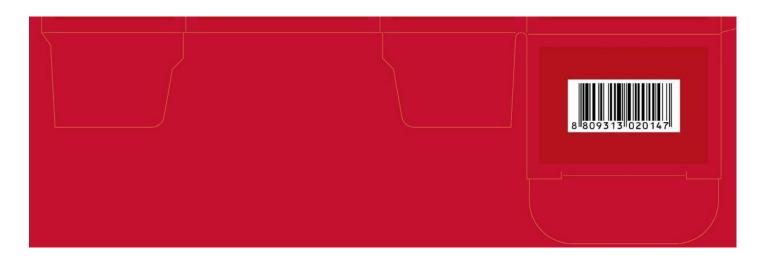
Indications and usage: 1. Tighten the lid after using it. 2. Don't keep it in the place where the temperature is extremely hot or low and exposed to the direct sunlight. 3. Use it in the evening after washing face.

DOSAGE AND ADMINISTRATION

Dosage and administration: Apply after washing face. Take a small amount of cream about 1.0-1.5g on to the fingertips and gently massage over the entire face. (cheeks, nostrils, corners, forehead, and the

PACKAGE LABEL. PRINCIPAL DISPLAY PANEL





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octinoxate, titanium dioxide, zinc oxide cream

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:60778-010
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
OCTINOXATE (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	OCTINOXATE	3 mg in 40 mL	
Titanium dioxide (UNII: 15FIX9 V2JP) (TITANIUM DIOXIDE - UNII:15FIX9 V2JP)	Titanium dioxide	2 mg in 40 mL	
Zinc oxide (UNII: SOI2LOH54Z) (ZINC CATION - UNII:13S1S8SF37)	ZINC CATION	2 mg in 40 mL	

Inactive Ingredients			
Ingredient Name	Strength		
Water (UNII: 059QF0KO0R)			
Butylene glycol (UNII: 3XUS85K0RA)			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:60778-010-01	40 mL in 1 CARTON		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part352	09/01/2013		

Labeler - Lycoris Co., Ltd. (688301521)

Registrant - Lycoris Co., Ltd. (688301521)

Establishment				
Name	Address	ID/FEI	Business Operations	
Lycoris Co., Ltd.		688301521	manufacture(60778-010)	

Revised: 10/2013 Lycoris Co., Ltd.